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L2	(angiotensin adj II) near10 receptor	1687	L2
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L1	(angiotensin adj II) near10 receptor	1563	L1

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Text Version	□ 1: Regul Pept. 1993 Mar 19;44(2):95-107.	Related Articles, Link			
Entrez PubMed Overview Help FAQ Tutorial	Immunohistochemical mapping of angiotensin AT1 receptors in the brain.				
New/Noteworthy E-Utilities	Phillips MI, Shen L, Richards EM, Raizada MK.				
PubMed Services	Department of Physiology, College of Medicine, Gainesville, FL 32610.				
Journals Database MeSH Database Single Citation Matcher Batch Citation Matcher Clinical Queries LinkOut Cubby	A new approach to study angiotensin receptor distribution in the brain has been taken by developing antibodies to partial sequence of the angiotensin II (AII) type-1 receptor subtype (AT1) and demonstrating the presence of receptors with immunohistochemical staining. The antibody to a portion of the 3rd cytoplasmic loop of the AT1 receptor revealed distinctive punctate immunoreactive staining on cell bodies. The cell bodies were distributed in the forebrain in persyentricular and suprapptic puclei, the organizm vasculosum lamina terminalis				
Related Resources Order Documents NLM Gateway	forebrain in paraventricular and supraoptic nuclei, the organum vasculosum lamina terminali median preoptic area and subfornical organ. In the brainstem, the entire locus coeruleus was stained, together with the adjacent mesencephalic and motor nuclei of the trigeminal nerve.				

PMID: 8469778 [PubMed - indexed for MEDLINE]

the cellular localization of AII receptors.

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The auditory system including the cochlear nucleus and superior olivary nuclei were stained.

nucleus ambiguous. Sites where AT2 receptors are located were not stained or staining was

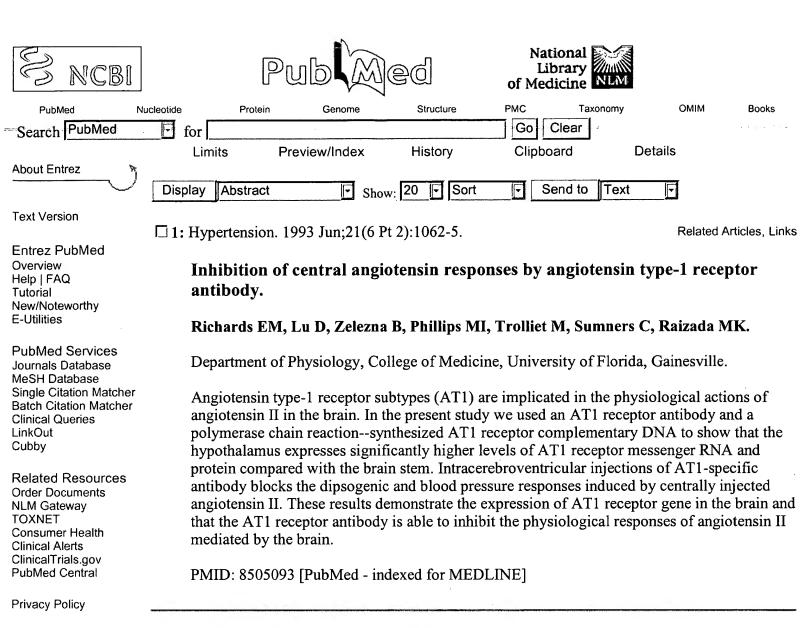
nucleus of the solitary tract, the 12th nerve nuclei, the rostroventral lateral area and the

limited to specific area such as the medial accessory nucleus of the inferior olive.

In the medulla, all the structures involved in blood pressure control were stained including the

Immunocytochemical staining of AT1 receptors provides a new and more precise approach to

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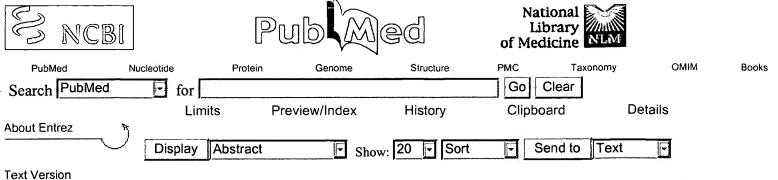
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□ 1: Am J Physiol. 1993 Jun;264(6 Pt 2):F989-95.

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Immunohistochemical localization of rat angiotensin II AT1 receptor.

Paxton WG, Runge M, Horaist C, Cohen C, Alexander RW, Bernstein KE.

Department of Pathology, Emory University, Atlanta, Georgia 30322.

To study receptors for angiotensin II, polyclonal rabbit anti-peptide antisera were prepared against the peptide QDDCPKAGRHC corresponding to amino acids 15-24 of the rat AT1A and AT1B receptors. Western analysis of rat tissues showed a major band of approximately 43 kDa. The antisera immunoprecipitated AT1-receptor protein produced in vitro. Immunohistochemical analysis of rat tissues showed intense staining of arterial and arteriolar smooth muscle. Other tissues that contained AT1-receptor protein included hepatocytes, the zona glomerulosa of the adrenal gland, and the smooth muscle of the bronchus, gut, ureter, and epididymis. In the kidney, intense staining was observed in all small arteries and arterioles. Both afferent and efferent arterioles contain approximately equal intensities of immunoreactive AT1 protein. The inner stripe of the outer medulla has a moderate level of receptors within thick ascending limb epithelium. Proximal tubular epithelium also expresses receptor protein. Glomerular immunoreactive AT1 protein is found within mesangial cells and varies in intensity among different rat strains. Lewis and Wistar rats demonstrated moderate glomerular staining, whereas the CD and Sprague-Dawley strains showed lesser levels of reactivity. The fact that glomerular mesangial cells are the primary locus of angiotensin II action within the glomerulus.

PMID: 7686719 [PubMed - indexed for MEDLINE]

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